



PATENT ABSTRACTS OF JAPAN

(11) Publication number: 06153947 A

(43) Date of publication of application: 03.06.1994

(51) Int. Cl. C12N 9/88
 A61K 37/56
 //(C12N 9/88, C12R 1:37)

(21) Application number: 05177458
 (22) Date of filing: 24.06.1993
 (30) Priority: 26.06.1992 JP 04192882

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(54) **CHONDROITINASE ABC, ITS PRODUCTION
 AND MEDICINE COMPOSITION**

(57) Abstract:

PURPOSE: To obtain a new enzyme useful as a therapeutic agent for herniated disk free from side effect because of absence of impurities, high purity, high specific activity and excellent stability, also useful as a reagent for research having high reproducibility.

CONSTITUTION: Purified chondroitinase ABC not containing endotoxin, having contents of nucleic acid and protease of a detection limit. Molecular weight:

100,000 Dalton. Isoelectric point: pH 8.2-pH 8.5. Optimum pH and optimum temperature: optimum pH 8-8.2 and optimum temperature 37°C in the case of chondroitin sulfate as a substrate in a tris-hydrochloric buffer solution. Influence by a metal; inhibition of activity with Zn^{2+} , Ni^{2+} , etc. End amino acids: Ala of N end and Pro of C end. Specific activity, $\geq 300V/mg$, etc. The enzyme is obtained by removing nucleic acid from an extracted solution of the enzyme yielded from a cell producing the enzyme and successively chromatographing the extracted solution with a weak cation exchange resin and a strong cation exchange resin.

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🔍 Title: **JP06153947A2: CHONDROITINASE ABC, ITS PRODUCTION AND MEDICINE COMPOSITION**

🔍 Derwent Title: Crystallisable, purified chondroitinase ABC - used in compsn. together with serum albumin and gelatin to treat intervertebral disc displacement
[\[Derwent Record\]](#)

🔍 Country: **JP Japan**

🔍 Kind: **A**

🔍 Inventor: **HASHIMOTO SHINICHI;
 MOCHIZUKI HIDEO;
 HAMAI AKIO;**

🔍 Assignee: **SEIKAGAKU KOGYO CO LTD**
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🔍 Published / Filed: **1994-06-03 / 1993-06-24**

🔍 Application Number: **JP1993000177458**

🔍 IPC Code: Advanced: **C12N 9/88;**
 Core: more...
 IPC-7: **A61K 37/56; C12N 9/88;**

🔍 Priority Number: **1992-06-26 JP1992000192882**

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CONSTITUTION: Purified chondroitinase ABC not containing endotoxin, having contents of nucleic acid and protease of a detection limit. Molecular weight: 100,000 Dalton. Isoelectric point: pH 8.2-pH 8.5. Optimum pH and optimum temperature: optimum pH 8-8.2 and optimum temperature 37°C in the case of chondroitin sulfate as a substrate in a tris-hydrochloric buffer solution. Influence by a metal; inhibition of activity with Zn²⁺, Ni²⁺, etc. End amino acids: Ala of N end and Pro of C end. Specific activity, ≥300U/mg, etc. The enzyme is obtained by removing nucleic acid from an extracted solution of the enzyme yielded from a cell producing the enzyme and successively chromatographing the extracted solution with a weak cation exchange resin and a strong cation exchange resin.

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